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## IN THE CLAIMS

Please cancel claims 1, 5-12 and 23-26. Please amend claims 2-4, 13-22 and 27-31. Claim 2 incorporates subject matter of claim 12. Claim 3 incorporates subject matter of claim 9.

## 1. (cancelled)

2. (presently amended) A method for site-specific incorporation of acyclonucleotides into DNA, comprising:

reacting an archaeon Family B DNA polymerase with a primed DNA template and nucleotide solution containing at least one acyclonucleotide to produce fragments of DNA with the acyclonucleotide covalently attached to the 3' terminal residue—;

wherein the DNA polymerase is encoded by an isolated DNA fragment that hybridizes in a Southern blot to an isolated DNA fragment selected from the group consisting of a DNA fragment having nucleotides 1-1274 of SEQ ID NO:4, a DNA fragment having nucleotides 291-1772 of SEQ ID NO:4, a DNA fragment having nucleotides 3387-3533 of SEQ ID NO:4, a DNA fragment having nucleotides 4704-5396 of SEQ ID NO:4, and a DNA fragment having nucleotides 4718-5437 of SEQ ID NO:4, wherein hybridization is conducted under the following conditions: a) hybridization: 0.75 M NaCl, 0.15 M Tris, 10 mM EDTA, 0.1% sodium pyrophosphate, 0.1% sodium lauryl sulfate, 0.03% BSA, 0.03% Ficoll 400, 0.03% PVP and 100 µg/ml boiled calf thymus DNA at 50°C for about 12 hours and; b) wash: 3X30 minutes with 0.1X SET, 0.1% SDS, 0.1% sodium pyrophosphate and 0.1 M phosphate buffer at 45°C.

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- 3. (presently amended) A method for site-specific incorporation of derivatized acyclonucleotides into DNA, comprising: reacting an archaeon Family B DNA polymerase with a primed DNA template and a nucleotide solution containing at least one derivatized acyclonucleotide to produce fragments of DNA with the derivatized acyclonucleotide covalently attached to the 3' terminal residue.; wherein the DNA polymerase has at least 30% primary amino acid sequence identity with Vent DNA polymerase.
- 4. (presently amended) The method of claims 1 2 or 3 wherein the derivative-acyclonucleotide comprises a detection reagent.

## 5-12 (cancelled)

- 13. (presently amended) The method of claims 1-3-2, wherein the DNA polymerase is selected from the group consisting of Vent, Deep Vent, Pfu and 9°N DNA polymerases.
- 14. (presently amended) The method of claims 1 3 2 or 3, wherein the DNA polymerase has been mutated by substitution of an amino acid residue at a site corresponding to A488, L492, A493 and Y499 in Vent polymerase.
- 15. (presently amended) The method of claims 1-3-2 or 3, wherein the DNA polymerase has been mutated by substitution of an amino acid residue corresponding to A488 in Vent polymerase with L, I, V, F, S or C.
- 16. (presently amended) The method of claims 1-3 2 or 3, wherein the DNA polymerase has been mutated by substitution of an amino acid residue corresponding to A488 in Vent DNA polymerase with L.

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- 17. (presently amended) The method of claims 1-3-2 or 3, wherein the DNA polymerase has been mutated by substitution of an amino acid residue corresponding to Y499 in Vent DNA polymerase with L.
- 18. (presently amended) The method of claims 1–3–2 or 3, wherein the DNA polymerase is a mutant selected from the group consisting of Vent (A488L), Vent' (Y499L) and 9°N (A485L) DNA polymerases.
- 19. (presently amended) The method of claims 2 or 3, wherein the acyclonucleotide is incorporated to an extent greater than that of a corresponding dideoxynucleotide.
- 20. (presently amended) The method of claims 2 or 3 wherein the acyclonucleotide is incorporated to an extent of at least, approximately, two-fold greater than incorporation of a corresponding dideoxynucleotide.
- 21. (presently amended) The method of claims 2 or 3 wherein the acyclonucleotide is incorporated to an extent at least, approximately, fivefold greater than incorporation of the corresponding dideoxynucleotide.
- 22. (presently amended) The method of claims 2 or 3 wherein the acyclonucleotide is incorporated to an extent at least, approximately, ninefold greater than incorporation of the corresponding dideoxynucleotide.

## 23-26 (cancelled)

27. (presently amended) The method of claims  $\frac{1-3}{2}$  wherein the DNA polymerase is additionally thermostable.

- 28. (presently amended) The method of claims 1-3  $\underline{2}$  wherein the DNA polymerase has no detectable exonuclease activity.
- 29. (presently amended) The method of claims 1-3 2 wherein the DNA polymerase has been mutated so as to have an exonuclease activity of less than about 5% of the exonuclease activity of the unmodified enzyme.
- 30. (presently amended) The method of claims 1-3-2 wherein the DNA polymerase has been mutated so as to have an exonuclease activity of less than about 25% of the exonuclease activity of the unmodified enzyme.
- 31. (presently amended) The method of claims 1-3-2 further comprising the step of employing the resulting sequence-specific termination product or products in DNA sequence determination.